

# Exhibit A

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Substitute for form 1449/PTO

**INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

Sheet 1 of 3

**Complete if Known**

Application Number	10/593,217
Filing Date	September 15, 2006
First Named Inventor	KING, et al.
Art Unit	Not Yet Known 1623
Examiner Name	Not Yet Known Ganapathy Krishnan
Attorney Docket Number	891-A-PCT-US

U. S. PATENT DOCUMENTS					
Examiner Initials*	Cite No. <sup>1</sup>	Document Number Number-Kind Code <sup>2</sup> (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
/G.K./	1	US- 4,175,200	11-20-1979	Hunter, et al.	
	2	US- 4,385,055	05-24-1983	Klayman, et al.	
	3	US- 4,447,427	05-08-1984	Klayman, et al.	
	4	US- 4,684,747	08-04-1987	Satorelli, et al.	
	5	US- 4,696,938	09-29-1987	Satorelli, et al.	
	6	US- 4,849,563	07-18-1989	Satorelli, et al.	
	7	US- 5,101,072	03-31-1992	Satorelli, et al.	
	8	US- 5,214,068	05-25-1993	Satorelli, et al.	
	9	US- 5,256,820	10-26-1993	Satorelli, et al.	
	10	US- 5,637,619	06-10-1997	Satorelli, et al.	
	11	US- 5,767,134	06-16-1998	Li, et al.	
	12	US- 6,040,338	03-21-2000	Satorelli, et al.	
/G.K./	13	US- 6,696,487	02-24-2004	Gerusz, et al.	
	14	US- 6,855,695	02-15-2005	Xu, et al.	
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FOREIGN PATENT DOCUMENTS						
Examiner Initials*	Cite No. <sup>1</sup>	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages Or Relevant Figures Appear	T <sup>6</sup>
		Country Code <sup>3</sup> Number <sup>4</sup> Kind Code <sup>5</sup> (if known)				
/G.K./	15	WO/2002/030424	04-18-2002	Doyle, et al.		

Examiner Signature	/Ganapathy Krishnan/	Date Considered	05/08/2008
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		Examiner Name	Not Yet Known		
Sheet	2	of	3	Attorney Docket Number	891-A-PCT-US

OTHER PRIOR ART—NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
/G.K./	16	Baumann RP, et al., 2004, "1,2-Bis(methylsulfonyl)-1-(2-chloroethyl)-2-[(methylamino)carbonyl]hydrazine (VNP40101M): II. Role of O6-alkylguanine-DNA alkyltransferase in cytotoxicity." Cancer Chemother Pharmacol. 53(4):288-295	
	17	Burchenal, et al., 1988, "Cancer: the Outlaw Cell," ed. Richard E Lafond, American Chemical Society: 204-205	
	18	Gura, et al., 1997, "Systems for identifying new drugs are often faulty," Science: 278(5340):1041-2	
	19	Lee, et al., 2002, "Toxicological evaluation of 1,2 bis("methylsulfonyl)-1-(2-chloroethyl)-2(methylaminocarbonyl) hydrazine (VNP40101M), novel alkylating Agent with Potential Antitumor Activity, with Intravenous Administration in Rats and Dogs", International Journal of Toxicology. Vol. 23: 23-39	
	20	Hrubiec, et al., 1986, "Synthesis and evaluation of 1-(arylsulfonyl)-2-[(methoxycarbonyl)sulfonyl]-1 methylhydrazines ++ + as antineoplastic agents." J Med Chem. 29(9):1777-9	
	21	Hrubiec, et al., 1986, "Synthesis and evaluation of 2-substituted 1-methyl-1-(4-tolylsulfonyl)hydrazines as antineoplastic agents," J Med Chem. 29(7):1299-301	
	22	Murren, et al., 2005, "A phase I and pharmacokinetic study of VNP40101M, a new alkylating agent, in patients with advanced or metastatic cancer", Investigational New Drugs. Vol 23: 123-135	
	23	Mao, et al., 2002, "Pharmacokinetics, Mass Balance, and Tissue Distribution of a Novel DNA Alkylating Agent, VNP40101M, in Rats", AAPS PharmSci: 4(4)24	
	24	Penketh, et al., 1994, "Studies on the mechanism of decomposition and structural factors affecting the aqueous stability of 1,2-bis(sulfonyl)-1-alkylhydrazines", J Med Chem 37: 2912-2917	
/G.K./	25	Penketh, et al., 2000, "Comparison of DNA lesions produced by tumor-inhibitory 1,2-bis(sulfonyl)hydrazines and chloroethylnitrosoureas", Biochem Pharmacol 59:283-91	

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Sheet	3	of	3	Attorney Docket Number	891-A-PCT-US

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/G.K./	26	Pratviel, et al., 1989, "Cytotoxic and DNA-damaging Effects of 1,2-bis(sulfonyl)hydrazines on Human Cells of the Mer+ and Mer- phenotype", Cancer Biochem Biophys 10:365-75 (abstract only)	
	27	Shyam, et al., 1985, "Synthesis and evaluation of N,N'-bis(arylsulfonyl)hydrazines as antineoplastic agents" J Med Chem 28:525-7	
	28	Shyam, et al., 1986, "1,2-bis(arylsulfonyl)hydrazines. 2. The influence of arylsulfonyl and alkylsulfonyl substituents on antitumor and alkylating activity", J Med Chem 29:1323-5	
	29	Shyam, et al., 1987, "1,2-Bis(sulfonyl)hydrazines. 3. Effects of structural modification on antineoplastic activity", J Med Chem 30:2157-61	
	30	Shyam, et al., 1990, "Synthesis and evaluation of 1,2,2-tris(sulfonyl)hydrazines as antineoplastic and trypanocidal agents", J Med Chem. 33(8):2259-64	
	31	Shyam, et al., 1993, "Synthesis and evaluation of 1-acyl-1,2-bis(methylsulfonyl)-2-(2-chloroethyl)hydrazines as antineoplastic agents", J Med Chem 36:3496-502	
	32	Shyam, et al., 1996, Antitumor 2-(aminocarbonyl)-1,2-bis(methylsulfonyl)-1-(2-chloroethyl)- hydrazines", J Med Chem 39:796-801	
	33	Giles, et al., 2004, "A Phase I and Pharmacokinetic Study of VNP40101M, a Novel Sulfonylhydrazine Alkylating Agent, in Patients with Refractory Leukemia.", Clinical Cancer Research, Vol. 10, Pages 2908-2917	
	34	Ishiguro, et al., 2005, "Role of O-alkylguanine-DNA alkyltransferase in the cytotoxic activity of cloreztazine", Mol Cancer Ther, Vol. 4 (11), Pages 1755-1763	
/G.K./	35	Rice, et al., 2005, "Differential inhibition of cellular glutathione reductase activity by isocyanates generated from the antitumor prodrugs Cloretazine and BCNU.", Biochemical Pharmacology, Vol. 69, Pages 1463-1472	

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